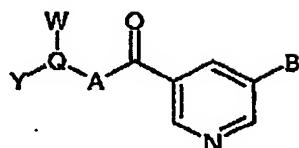


## CLAIMS:-

## 1 A compound of the general formula



5 or pharmaceutically acceptable prodrugs, salts, hydrates, solvates, crystal forms or diastereomers therof, wherein:

A is selected from O, S, NR1, where R1 is selected from H, C<sub>1-4</sub> alkyl;

10 B is aryl, hetaryl optionally substituted with 0-3 substituents independently chosen from halogen, C<sub>1-4</sub> alkyl, CF<sub>3</sub>, CN, aryl, hetaryl, OH, OCF<sub>3</sub>, OC<sub>1-4</sub>alkyl, OC<sub>2-6</sub>alkylNR2R3, Oaryl, Ohetaryl, CO<sub>2</sub>R2, CONR2R3, NR2R3, C<sub>1-4</sub> alkylNR2R3, NR4C<sub>1-4</sub>alkylNR2R3, NR2COR3, OC(O)NR2R3, NR4CONR2R3, NR2SO<sub>2</sub>R3; and R2, R3 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl heterocycl, aryl, hetaryl, C<sub>1-4</sub>alkyl aryl, C<sub>1-4</sub> alkyl hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR5; and R4 is selected from H, C<sub>1-4</sub> alkyl; and R5 is selected from H, C<sub>1-4</sub> alkyl;

15 Q is a bond, or C<sub>1-4</sub> alkyl;

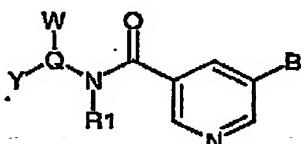
20 W is selected from H, C<sub>1-4</sub>alkyl, C<sub>2-6</sub>alkenyl; where C<sub>1-4</sub>alkyl or C<sub>2-6</sub>alkenyl may be optionally substituted with C<sub>1-4</sub>alkyl, OH, OC<sub>1-4</sub>alkyl, NR<sub>6</sub>C(O)R7, CONR6R7, OR6, NR6R7; and R6, and R7 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl cycloalkyl, C<sub>1-4</sub> alkyl heterocycl, aryl, hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR8 and R8 is selected from H, C<sub>1-4</sub> alkyl;

25 Y is H, aryl or hetaryl optionally substituted with 0-3 substituents independently chosen from halogen, C<sub>1-4</sub> alkyl, CF<sub>3</sub>, aryl, hetaryl, OH, OCF<sub>3</sub>, CN, C<sub>2-4</sub> alkynyl, OC<sub>1-4</sub>alkyl, OC<sub>2-6</sub>alkylNR9R10, Oaryl, Ohetaryl, CO<sub>2</sub>R9, CONR9R10, NR9R10, C<sub>1-4</sub> alkylNR9R10,

NR11C<sub>1-4</sub>alkylNR9R10, NR9COR10, NR11CONR9R10, NR9SO<sub>2</sub>R10; and R9, R10 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl heterocycll, aryl, hetaryl, C<sub>1-4</sub> alkyl aryl, C<sub>1-4</sub> alkyl hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR12; and R11 is selected from H, C<sub>1-4</sub> alkyl; and

5 R12 is selected from H, C<sub>1-4</sub> alkyl.

2. A compound according to claim 1 of the general formula II:



10 or pharmaceutically acceptable prodrugs, salts, hydrates, solvates, crystal forms or diastereomers thereof, wherein:

R1 is selected from H, C<sub>1-4</sub> alkyl;

15 B is aryl, hetaryl optionally substituted with 0-3 substituents independently chosen from halogen, C<sub>1-4</sub> alkyl, CF<sub>3</sub>, aryl, hetaryl, OH, OCF<sub>3</sub>, OC<sub>1-4</sub>alkyl, OC<sub>2-5</sub>alkylNR2R3, Oaryl, Ohetaryl, CO<sub>2</sub>R2, CONR2R3, NR2R3, C<sub>1-4</sub> alkylNR2R3, NR4C<sub>1-4</sub>alkylNR2R3, NR2COR3, NR4CONR2R3, NR2SO<sub>2</sub>R3; and R2, R3 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl heterocycll, aryl, hetaryl, C<sub>1-4</sub> alkyl aryl, C<sub>1-4</sub> alkyl hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR5; and R4 is selected from H, C<sub>1-4</sub> alkyl; and R5 is selected from H, C<sub>1-4</sub> alkyl;

20 Q is a bond, or C<sub>1-4</sub> alkyl;

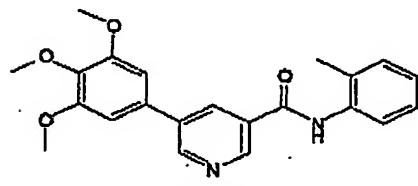
25 W is selected from H, C<sub>1-4</sub>alkyl, C<sub>2-6</sub>alkenyl; where C<sub>1-4</sub>alkyl or C<sub>2-6</sub>alkenyl may be optionally substituted with C<sub>1-4</sub>alkyl, OH, OC<sub>1-4</sub>alkyl, NR6R7; and R6, and R7 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl cycloalkyl, C<sub>1-4</sub> alkyl heterocycll, aryl, hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR8 and R8 is selected from H, C<sub>1-4</sub> alkyl;

Y is H, aryl or hetaryl optionally substituted with 0-3 substituents independently chosen from halogen, C<sub>1-4</sub> alkyl, CF<sub>3</sub>, aryl, hetaryl, OH, OCF<sub>3</sub>, OC<sub>1-4</sub>alkyl, OC<sub>2-5</sub>alkylNR9R10, Oaryl, Ohetaryl, CO<sub>2</sub>R9, CONR9R10, NR9R10, C<sub>1-4</sub>alkylNR9R10, NR11C<sub>1-4</sub>alkylNR9R10, NR9COR10, NR11CONR9R10, NR9SO<sub>2</sub>R10; and R9, R10 are each independently H, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkyl heterocycl, aryl, hetaryl, C<sub>1-4</sub>alkyl aryl, C<sub>1-4</sub> alkyl hetaryl, or may be joined to form an optionally substituted 3-8 membered ring optionally containing an atom selected from O, S, NR12; and R11 is selected from H, C<sub>1-4</sub> alkyl; and R12 is selected from H, C<sub>1-4</sub> alkyl.

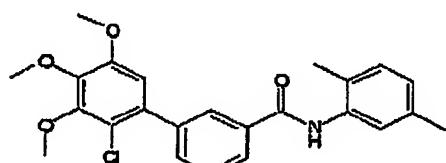
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3. A compound according to claim 1 wherein the compound is selected from the group consisting of:

10

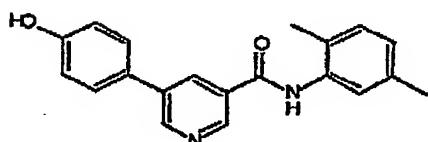


C22H22N2O4

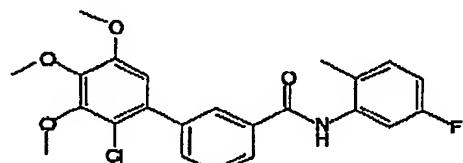


C23H23ClN2O4

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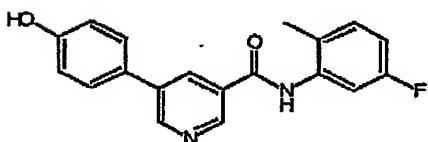


C20H18N2O2

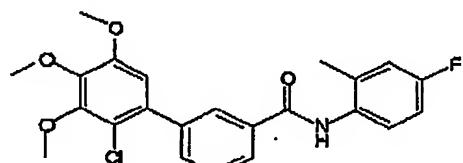


C22H20ClFN2O4

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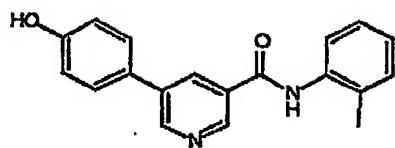


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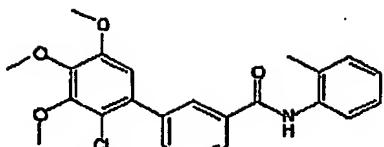


C22H20ClFN2O4

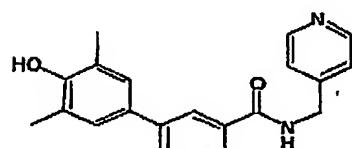
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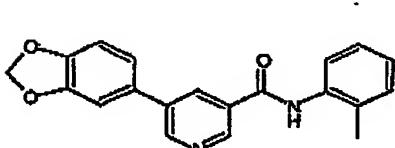
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C22H21ClN2O4

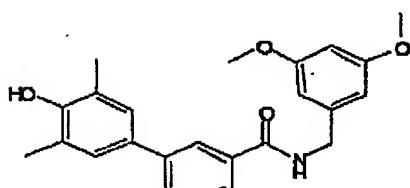


C20H18N3O2

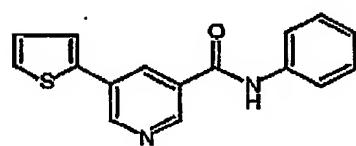


C20H16N2O3

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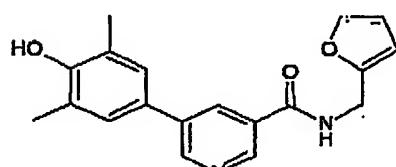


C23H24N2O4

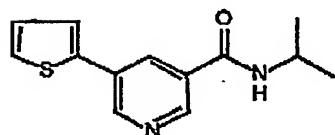


C16H12N2OS

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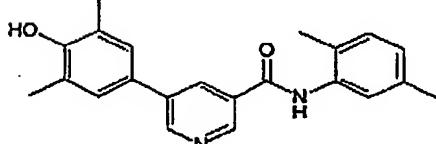


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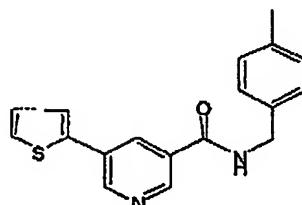


C13H14N2OS

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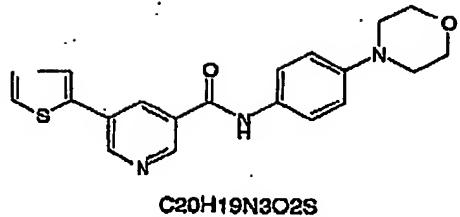
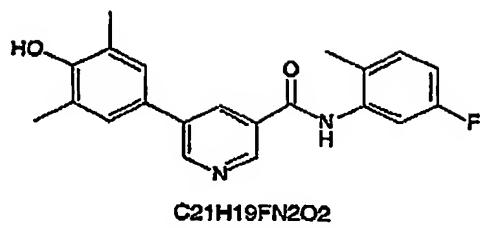


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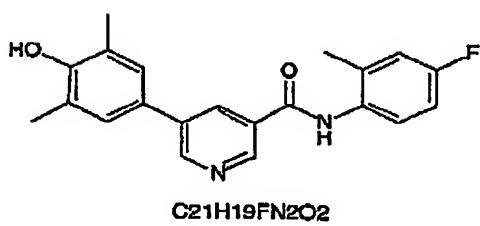


C18H16N2OS

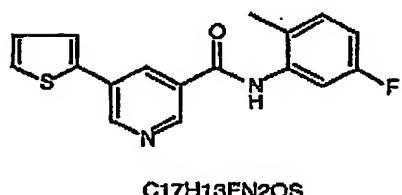
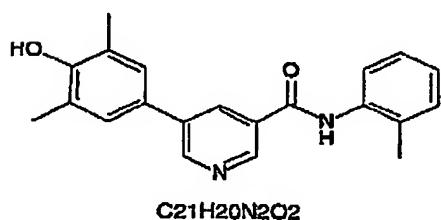
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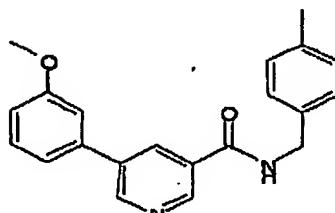
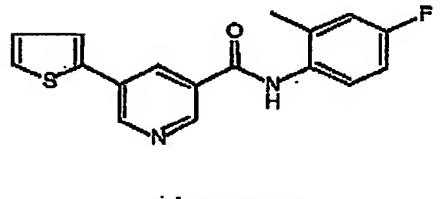
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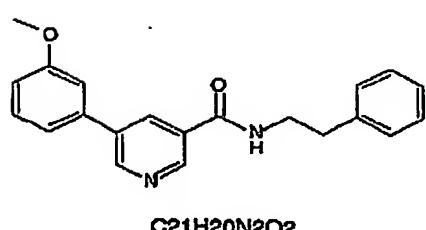
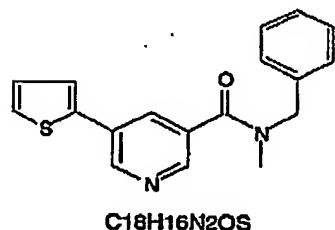
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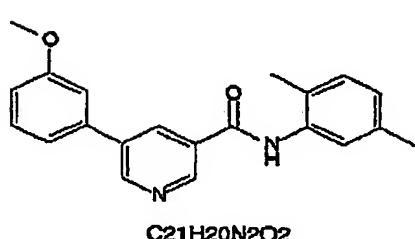
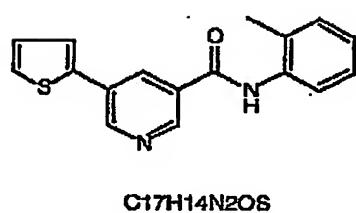
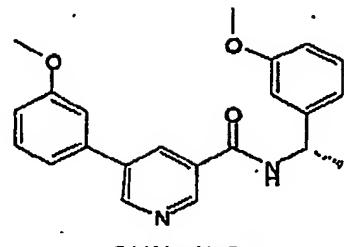
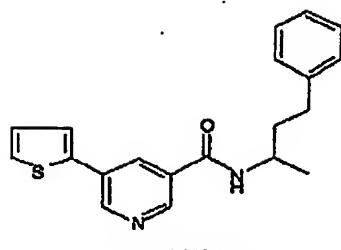
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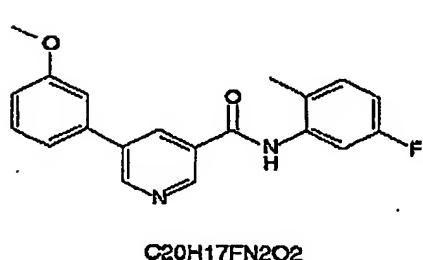
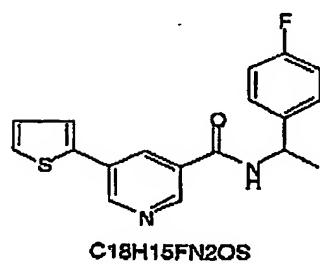
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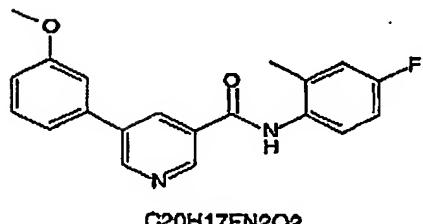
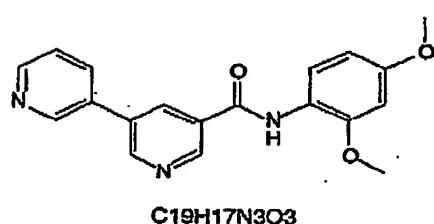
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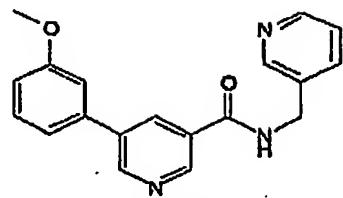
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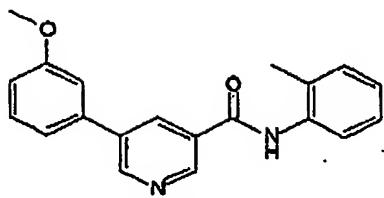
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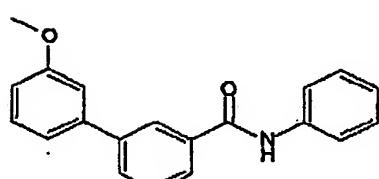


C19H17N3O2

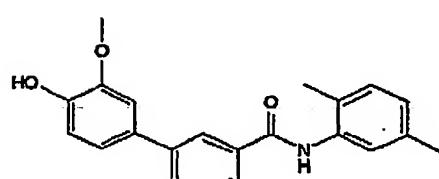


C20H18N2O2

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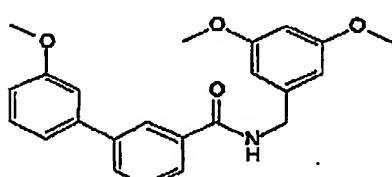


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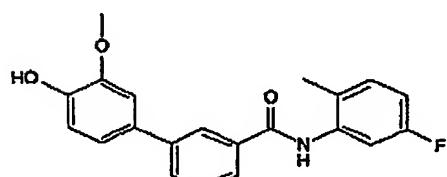


C21H20N2O3

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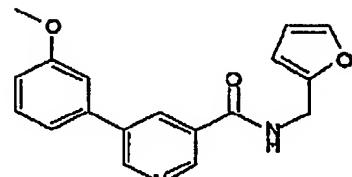


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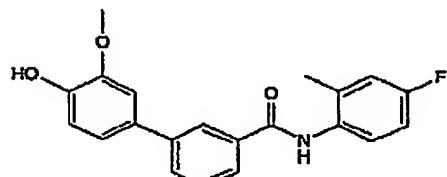


C20H17FN2O3

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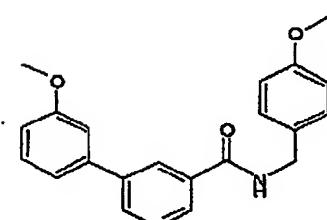


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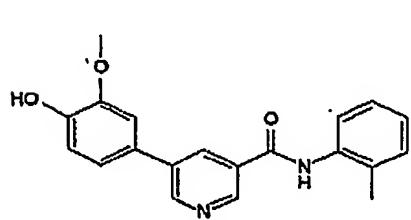


C20H17FN2O3

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C21H20N2O3



C20H18N2O3

4. A composition comprising a carrier and at least one compound of any one of claims 1 to 3.
5. A composition comprising a carrier and at least one compound of any one of claims 1 to 4.
- 5 6. A method of treating a tyrosine kinase-associated disease state in a subject, the method comprising administering a therapeutically effective amount of at least one compound of any one of claims 1 to 4 or a therapeutically effective amount of a composition of claim 5.
7. A method according to claim 6 wherein the disease state is selected from the group consisting of Atopy, such as Allergic Asthma, Atopic Dermatitis (Eczema), and Allergic Rhinitis; Cell Mediated Hypersensitivity, such as Allergic Contact Dermatitis and Hypersensitivity Pneumonitis; Rheumatic Diseases, such as Systemic Lupus Erythematosus (SLE), Rheumatoid Arthritis, Juvenile Arthritis, Sjögren's Syndrome, Scleroderma, Polymyositis, Ankylosing Spondylitis, Psoriatic Arthritis; Other autoimmune diseases such as Type I diabetes, autoimmune thyroid disorders, and Alzheimer's disease; Viral Diseases, such as Epstein Barr Virus (EBV), Hepatitis B, Hepatitis C, HIV, HTLV 1, Varicella-Zoster Virus (VZV), Human Papilloma Virus (HPV); Cancer, such as fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, 10 endothelioma, lymphangiosarcoma, lymphangioendothelioma, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary 15 adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilms' tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, 20 pinealoma, hemangioblastoma, acoustic neuroma, oligodendrogloma, meningioma, melanoma, neuroblastoma, and retinoblastoma, and carcinomas forming from tissue of the breast, prostate, kidney, bladder or colon, and neoplastic 25 30

disorders arising in adipose tissue, such as adipose cell tumors, e.g., lipomas, fibrolipomas, lipoblastomas, lipomatosis, hibernomas, hemangiomas and/or liposarcomas.

5        8. The use of at least one of the compounds of any one of claims 1 to 4 in the preparation of a medicament for the treatment of a tyrosine kinase-associated disease state.

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